

**HRT Medication Adherence****Data File:** ht_adh_ep_fu_pub **File Date:** 07/22/2005 **Structure:** Multiple rows per participant **Population:** E+P participants

Participant ID**Variable #** 1**Usage Notes:** none**Sas Name:** ID**Categories:** Study: Administration**Sas Label:** Participant ID**Type:** Continuous

Visit type for HRT adherence period**Variable #** 2**Usage Notes:** none**Sas Name:** ADHVTYP**Categories:** Study Interventions: HRT Intervention/Management**Sas Label:** Visit type for HRT adherence period**Type:** Categorical**Values**

3	Annual Visit
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Visit year for HRT adherence period**Variable #** 3**Usage Notes:** none**Sas Name:** ADHVYR**Categories:** Study Interventions: HRT Intervention/Management**Sas Label:** Visit year for HRT adherence period**Type:** Continuous

Days from rand to start of HRT adherence period

Days from randomization to start of HRT adherence period.

Variable # 4**Usage Notes:** none**Sas Name:** STARTDY**Categories:** Study Interventions: HRT Intervention/Management**Sas Label:** Days from rand to start of HRT adherence period**Type:** Continuous

Days from randomization to end of adherence period

Days from randomization to end of HRT adherence period.

Variable # 5**Usage Notes:** none**Sas Name:** ENDDY**Categories:** Study Interventions: HRT Intervention/Management**Sas Label:** Days from rand to end of HRT adherence period**Type:** Continuous

HRT medication adherence rate for the period**Variable #** 6**Usage Notes:** none**Sas Name:** ADHRATE**Categories:** Study Interventions: HRT Intervention/Management**Sas Label:** HRT medication adherence rate for the period**Type:** Continuous

**Was adherence collection performed during period**

Were HRT pill bottles collected to allow adherence determination for period? If not, no adherence rate can be calculated.

Variable # 7**Usage Notes:** none**Sas Name:** COLLECT**Sas Label:** Was adherence collection performed during period**Categories:** Study Interventions: HRT Intervention/Management**Type:** Categorical**Values**

0	No
1	Yes

Participant inactive in intervention during period

Was the participant inactive in the HRT intervention (i.e. not taking study pills) for all or part of the period.

Variable # 8**Usage Notes:** none**Sas Name:** STOPHRT**Sas Label:** Participant inactive in HRT interventin during period**Categories:** Study Interventions: HRT Intervention/Management**Type:** Categorical**Values**

0	No
1	Yes

Participant resumed HRT intervention during period

Did the participant resume HRT intervention (start taking study pills) during this period after having stopped?

Variable # 9**Usage Notes:** none**Sas Name:** RESUMEHRT**Sas Label:** Participant resumed HRT intervention during period**Categories:** Study Interventions: HRT Intervention/Management**Type:** Categorical**Values**

0	No
1	Yes

Participant lost-to-follow-up during period

Did the participant have a status of lost-to-follow-up during all or part of this HRT adherence period?

Variable # 10**Usage Notes:** none**Sas Name:** LOST**Sas Label:** Participant lost-to-follow-up during period**Categories:** Study Interventions: HRT Intervention/Management**Type:** Categorical**Values**

0	No
1	Yes



WHI Follow-Up Dataset
HRT Medication Adherence

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Participant deceased during period

Did the participant have a status of deceased during all or part of this HRT adherence period?

Variable # 11 **Usage Notes:** none
Sas Name: DEAD **Categories:** Study Interventions: HRT Intervention/Management
Sas Label: Participant deceased during period
Type: Categorical

Values

0	No
1	Yes

Were open label HRT meds dispensed during period

Were any open label HRT medications dispensed to the participant during this period?

Variable # 12 **Usage Notes:** Open label medications are not factored into the adherence rate calculation.
Sas Name: OPENLABEL **Categories:** Study Interventions: HRT Intervention/Management
Sas Label: Were open label HRT meds dispensed during period
Type: Categorical

Values

0	No
1	Yes

Participant switched from E-alone to E+P in period

Participant was switched from the unopposed estrogen study group to the estrogen+progesterone study group during this period (January 1995), due to PEPI trial results indicating long-term adherence to estrogen was not feasible in women with a uterus.

Variable # 13 **Usage Notes:** none
Sas Name: ERT2PERT **Categories:** Study Interventions: HRT Intervention/Management
Sas Label: Participant was switched from E-alone to E+P in period
Type: Categorical

Values

0	No
1	Yes